



DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration  
Rockville MD 20857

MAR 23 1988

Re: Novantrone  
Docket No. 88E-0067

The Honorable Donald J. Quigg  
Assistant Secretary of Commerce  
and Commissioner of Patents and Trademarks  
Washington, D.C. 20231

Dear Commissioner Quigg:

This is in regard to the application for patent extension for U.S. Patent No. 4,278,689, filed by the American Cyanamid Company, under 35 U.S.C. 156 et seq. We have reviewed the dates contained in the application and have determined the regulatory review period for Novantrone, the human drug claimed by the patent.

The total length of the review period for Novantrone is 3,145 days. Of this time, 1,836 days occurred during the testing phase and 1,309 days occurred during the approval phase. The periods of time were derived from the following dates:

1. The date an exemption under subsection 505 (i) of the Federal Food, Drug, and Cosmetic Act involving this drug product became effective: May 16, 1979.

The applicant claims April 16, 1979 as the date the notice of claimed investigational exemption (IND) for the drug became effective. However, FDA records indicate that the IND did not become effective until May 16, 1979.

2. The date the application was initially submitted with respect to the human drug product under subsection 505 (b) of the Federal Food, Drug, and Cosmetic Act: May 24, 1984.

The applicant claims a new drug application (NDA 19-297) was initially submitted on May 18, 1984. However, FDA did not receive the application until May 24, 1984.

3. The date the application was approved: December 31, 1987.

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U.S. TRADE  
COMMISSIONER

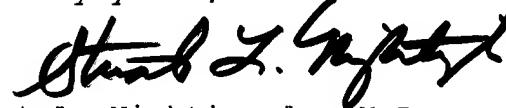
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FDA has verified the applicant's claim that NDA 19-297 was approved on December 31, 1987.

This determination of the regulatory review period by FDA does not take into account the effective date of the patent nor does it exclude one-half of the testing phase as required by 35 U.S.C. 156 (c)(2).

Please let us know if we can be of further assistance.

Sincerely yours,



Stuart L. Nightingale, M.D.  
Associate Commissioner  
for Health Affairs

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